

## RESEARCH ARTICLE

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## Efficient narrowband green OLEDs with TADF sensitizers combining multiple charge-transfer pathways†

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Multiple resonance (MR) thermally activated delayed fluorescence (TADF) emitters have attracted much attention for their narrow emission band and high efficiency. However, organic light-emitting devices based on MR-TADF emitters generally suffer from severe efficiency roll-off at high current density due to their relatively large singlet–triplet splitting energy. In this work, three TADF molecules are designed by an *ortho*-connected oligo carbazole donor and diphenyltriazine acceptor with the purpose of combining both through-bond charge transfer (TBCT) and through-space charge transfer (TSCT). Through fine tuning the proportion of intramolecular TBCT and TSCT by changing the number of carbazole units, a relatively high reverse intersystem crossing rate ( $k_{\text{RISC}}$ ) approaching  $10^6 \text{ s}^{-1}$  and radiative transition rate ( $k_{\text{F}}$ ) of over  $10^6 \text{ s}^{-1}$  are achieved. As a result, the devices using these molecules as emitters give maximum external quantum efficiency (EQE) of over 20% with extremely low efficiency roll-off of 9.9% at the practical luminance of  $1000 \text{ cd m}^{-2}$ . Furthermore, by using these materials as assistant hosts and sensitizers and DMAC-BN as a terminal emitter, high efficiencies of 23.9% and  $66.0 \text{ lm W}^{-1}$  are achieved with a full width at half maximum of 46 nm, featuring slow efficiency roll-off with EQE of 17.9% at the luminance of  $1000 \text{ cd m}^{-2}$ .

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## Introduction

Thermally activated delayed fluorescence (TADF) materials have attracted global attention from the academic and industrial communities since the pioneering work of the purely organic TADF organic light-emitting diodes (OLEDs) developed by Adachi and coworkers.<sup>1–3</sup> Owing to the unity exciton utilization comparable to noble-metal-containing phosphorescence materials and much higher than traditional fluorescence materials, as well as facile property modulation through manipulating the charge transfer (CT) characteristics, TADF materials hold great potential to achieve high performance OLEDs.<sup>4–6</sup> Through-bond charge transfer (TBCT) and through-space charge transfer (TSCT) are two general CT characteristics for developing TADF materials.<sup>4,6–9</sup> High radiative transition rate ( $k_{\text{F}}$ ) and reverse intersystem crossing (RISC) rate ( $k_{\text{RISC}}$ ) are indispensable for TADF materials to achieve high external quantum efficiency (EQE) and low efficiency

roll-off.<sup>10,11</sup> Generally, a considerable overlap of highest occupied molecular orbital (HOMO) and lowest unoccupied molecular orbital (LUMO) is conducive to a high oscillator strength ( $f$ ) and then a high  $k_{\text{F}}$ .<sup>12–14</sup> However, for achieving a high  $k_{\text{RISC}}$ , the most effective strategy is to reduce the splitting energy between the lowest triplet state ( $T_1$ ) and the lowest singlet state ( $S_1$ ), *i.e.*,  $\Delta E_{\text{ST}}$ , by decreasing the overlap of the HOMO and LUMO.<sup>1,4,13</sup> It follows that achieving high  $k_{\text{F}}$  and  $k_{\text{RISC}}$  simultaneously is of some difficulty in TADF materials, which is an obstacle for achieving high EQE and low efficiency roll-off simultaneously.

With regard to the TBCT-based TADF materials, appropriate donors/acceptors that possess suitable electron donating/accepting abilities are generally combined through a conjugated unit, and the torsion angles between the donors and acceptors should be subtly manipulated to give suitable HOMO–LUMO overlaps and thus high  $k_{\text{F}}$ .<sup>2,4,15–17</sup> As for the TSCT-based TADF materials, the CT process occurs through the spatial  $\pi$ – $\pi$  interaction of the donors and acceptors. The tiny overlaps of the HOMOs and LUMOs for this kind of material generally give high  $k_{\text{RISC}}$ .<sup>18–21</sup> It appears that concurrent intramolecular TBCT and TSCT might be an effective way to achieve high  $k_{\text{F}}$  and  $k_{\text{RISC}}$  simultaneously. Unfortunately, the majority of the reported intramolecular TSCT-based materials mainly emphasized the

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existence of TSCT without too much consideration of TBCT.<sup>20–22</sup> Although a few research studies have paid attention to the coexistence of intramolecular TBCT and TSCT,<sup>5,14,23,24</sup> systematic manipulation of the intramolecular TBCT and TSCT was not mentioned.

Due to the strong CT characteristic, D–A-type TADF emitters, including TBCT and TSCT type molecules, exhibit a broad electroluminescence (EL) band with a typical full-width at half-maximum (FWHM) of 80–100 nm. To address this crucial flaw, Hatakeyama *et al.*<sup>25,26</sup> reported a new design strategy featuring an alternating HOMO/LUMO pattern, based on the multiple resonance (MR) effect of boron and nitrogen/oxygen atoms. Most MR-based molecules possess a narrow emission band owing to the suppressed vibronic coupling between the ground ( $S_0$ ) and singlet excited ( $S_1$ ) states. Even though these MR-TADF emitters rendered excellent performance in OLED devices, severe efficiency roll-off at high current density can be observed due to their relatively large  $\Delta E_{ST}$ .<sup>25,27–30</sup> To overcome this drawback, great efforts have been made in molecular design and device design, such as increasing  $k_{RISC}$  and using a phosphor or TADF molecule as an assistant host (sensitizer) to improve the device performance and reduce the efficiency roll-off of the devices with MR-TADF materials as terminal emitters.<sup>5,31–37</sup>

In this research, three TADF molecules, *o*-Cz-TRZ, *o*-DCz-TRZ and *o*-TCz-TRZ, are constructed by an oligo carbazole donor and diphenyltriazine acceptor *ortho*-connected to a benzene linker (Fig. 1). The *ortho*-linkage between the donor and acceptor is conducive to inducing intramolecular TSCT characteristics for the short spatial distance of the donor and acceptor while retaining TBCT characteristics through the HOMO–LUMO overlap localized at the benzene linker. The proportions of intramolecular TBCT and TSCT are easily modulated by the number of carbazole units. Benefitting from the concurrent intramolecular TBCT and TSCT,  $k_F$  exceeding  $10^6 \text{ s}^{-1}$  and  $k_{RISC}$  approaching  $10^6 \text{ s}^{-1}$  are achieved simultaneously for all the developed materials.

*o*-DCz-TRZ with 70.0% intramolecular TSCT characteristic and 25.5% TBCT characteristic gives the highest  $k_{RISC}$  and photoluminescence quantum yield (PLQY) and a relatively high  $k_F$  among the three compounds, resulting in a maximum EQE of 20.3% and a low efficiency roll-off of 11.8% at  $1000 \text{ cd m}^{-2}$  for the device based on *o*-DCz-TRZ as the emitter. Furthermore, by using these materials as assistant hosts and sensitizers and DMAC-BN as a terminal emitter,<sup>29,30</sup> high efficiencies of 23.9% and  $66.0 \text{ lm W}^{-1}$  are achieved for the device sensitized by *o*-TCz-TRZ, featuring a FWHM of 46 nm and a slow efficiency roll-off with EQE of 17.9% at the luminance of  $1000 \text{ cd m}^{-2}$ . The current work provides a simple molecular design strategy of tunable intramolecular TBCT and TSCT for TADF molecules with simultaneous high  $k_F$  and  $k_{RISC}$  toward high-efficiency and low efficiency roll-off OLEDs.

## Results and discussion

### Synthesis and characterization

The molecular structures and synthetic routes are illustrated in Fig. 1 and Scheme S1 (ESI<sup>†</sup>). The intermediate 2-(2-fluorophenyl)-4,6-diphenyl-1,3,5-triazine and target product *o*-Cz-TRZ were synthesized according to the reported literature.<sup>38,39</sup> The chemical structures of all the materials were determined by <sup>1</sup>H NMR, <sup>13</sup>C NMR and mass spectra. Thermogravimetric analysis (Fig. S1a, ESI<sup>†</sup>) reveals that all the compounds exhibit good thermal stability with high decomposition temperatures ( $T_d$ , corresponding to a weight loss of 5 wt%) over  $360 \text{ }^\circ\text{C}$  (Table S1, ESI<sup>†</sup>). According to the differential scanning calorimetry (DSC) analysis, all the compounds exhibit obvious glass transition processes with glass transition temperatures ( $T_g$ ) of 62, 120 and  $165 \text{ }^\circ\text{C}$  for *o*-Cz-TRZ, *o*-DCz-TRZ and *o*-TCz-TRZ, respectively. No obvious melting processes were observed for *o*-DCz-TRZ and *o*-TCz-TRZ in the range of  $0\text{--}300 \text{ }^\circ\text{C}$  whereas *o*-Cz-TRZ exhibits a melting point ( $T_m$ ) of  $210 \text{ }^\circ\text{C}$ . In addition, distinct

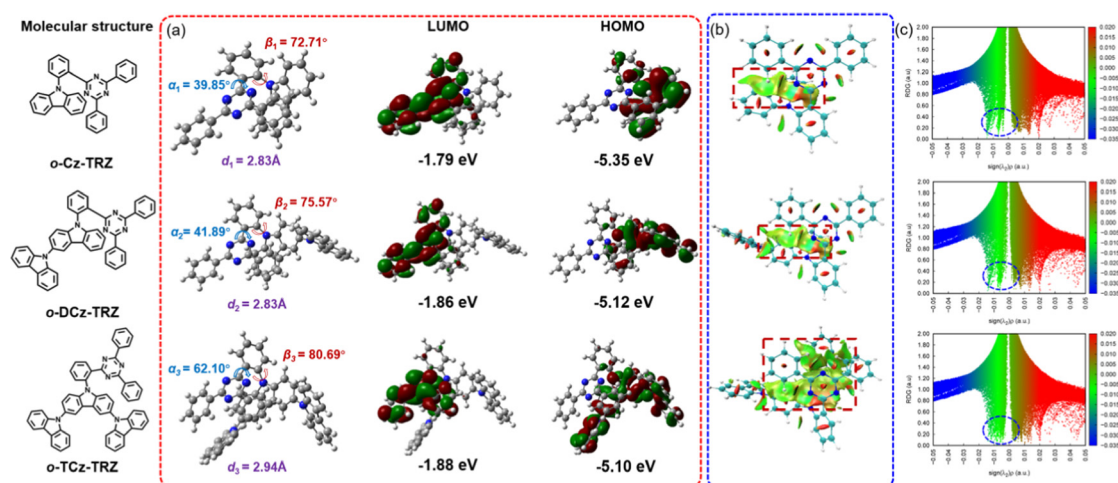


Fig. 1 Chemical structures and optimized molecular geometries of the ground states and FMO distributions and energy levels (a) simulated at the B3LYP/6-31G\*\* level. RDG isosurfaces (b) and RDG scattering diagrams (c) based on DFT simulations for *o*-Cz-TRZ, *o*-DCz-TRZ and *o*-TCz-TRZ, respectively.

non-equilibrium crystal transformation processes were observed for all the compounds (Fig. S1b, ESI<sup>†</sup>). Electrochemical properties were investigated by cyclic voltammetry (CV) measurements (Fig. S2, ESI<sup>†</sup>), and the HOMO energy levels were estimated to be  $-5.52$ ,  $-5.38$  and  $-5.37$  eV for *o*-Cz-TRZ, *o*-DCz-TRZ and *o*-TCz-TRZ, respectively, from the onset potentials of the redox peaks according to Eqn S1 (Table S1, ESI<sup>†</sup>).

### Computational simulations

To gain insight into the molecular structures and electron distributions, density functional theory (DFT) and time-dependent DFT (TD-DFT) simulations were conducted based on the B3LYP/6-31G\*\* level. As depicted in Fig. 1 in the ground state geometry, the *ortho*-connected donor and acceptor lead to approximately face-to-face alignment of the oligo carbazole and triazine rings on the phenylene bridge, which ensures the occurrence of intramolecular TSCT from the donor to the acceptor directly. Attributed to the *ortho*-linkage and the large steric hindrance, these molecules exhibit a twisted structure with torsion angles between the oligo carbazole donor and benzene linker ( $\beta$ , red curved arrow) of  $72.71^\circ$ ,  $75.57^\circ$ , and  $80.69^\circ$  for *o*-Cz-TRZ, *o*-DCz-TRZ and *o*-TCz-TRZ, respectively. Such large torsion angles are beneficial to separate the HOMOs and LUMOs, leading to small computational  $\Delta E_{\text{STS}}$  of 90, 50 and 30 meV for *o*-Cz-TRZ, *o*-DCz-TRZ and *o*-TCz-TRZ, respectively (Fig. 1). With increasing number of carbazole units, both the torsion angles of  $\alpha$  (the torsion angle between the benzene linker and diphenyltriazine acceptor) and  $\beta$  increase in varying degrees. Furthermore, the HOMO–LUMO overlap at the benzene linker is reduced, leading to low oscillator strength ( $f = 0.0052$ ,  $0.0040$  and  $0.0022$  for *o*-Cz-TRZ, *o*-DCz-TRZ and *o*-TCz-TRZ, respectively, as shown in Fig. S3, ESI<sup>†</sup>).

In order to investigate the intramolecular weak interactions of these materials, reduced density gradient (RDG) isosurfaces and scattering diagrams were performed in virtue of Multiwfn and visual molecular dynamics (VMD) (Fig. 1).<sup>40–42</sup> As illustrated in Fig. 1b, a noticeable non-covalent weak interaction between the oligo carbazole donor and diphenyltriazine acceptor can be observed in the RDG isosurfaces (the green area in the red dotted box), which demonstrates the existence of intramolecular TSCT

intuitively.<sup>5,7,19,21</sup> The non-covalent weak interactions are strengthened with increasing number of carbazole units, which can be embodied in the larger areas in the RDG isosurfaces. Besides, the more intensive spikes are observed as shown in the RDG scattering diagrams (Fig. 1c, the green spikes in the blue dotted circles), demonstrating the effectiveness of intramolecular TSCT modulation by the structure of the donor. Moreover, the interfragment charge transfer (IFCT) analysis based on the TD-DFT calculation results was conducted, which quantitatively depicted the contribution of electron transfer among the fragments (donor, acceptor and linker).<sup>5,19,21</sup> The interaction ratios of TSCT in the  $S_1$  state were calculated to be 0.6376, 0.7002 and 0.924 for *o*-Cz-TRZ, *o*-DCz-TRZ and *o*-TCz-TRZ, respectively, according to the reported literature.<sup>5</sup>

### Photophysical properties

Ultraviolet-visible (UV-vis) absorption and photoluminescence (PL) spectra of these molecules were measured in diluted toluene solution (Fig. 2a). For all the target molecules, the relatively strong absorption bands at around 310–350 nm mainly originate from the local excited (LE) state transition of the carbazole-based donors, and the tail absorption bands at around 350–440 nm could be attributed to the intramolecular charge-transfer (ICT) transition (Fig. 3a). The optical bandgaps ( $E_{\text{g}}$ s) are estimated to be 2.63, 2.52 and 2.52 eV for *o*-Cz-TRZ, *o*-DCz-TRZ and *o*-TCz-TRZ, respectively, from the onset of the absorption spectra. In the PL spectra, *o*-Cz-TRZ exhibits a blue emission peak at 460 nm while *o*-DCz-TRZ and *o*-TCz-TRZ exhibit sky-blue emission peaks at 495 and 492 nm, respectively. In comparison with *o*-Cz-TRZ, the bathochromic shift of the PL spectra for *o*-DCz-TRZ and *o*-TCz-TRZ should be attributed to the stronger electron-donating ability of the DCz and TCz donor units, which is consistent with the tendency of HOMO energy levels attained from the CV measurements. Nearly superimposed low-temperature fluorescence and phosphorescence spectra were observed in toluene solution (Fig. S5, ESI<sup>†</sup>), resulting in tiny  $\Delta E_{\text{STS}}$  approaching zero for all the materials (Table 1), which are consistent with the TD-DFT calculations. In accordance with the natural transition orbital (NTO) analysis of the  $S_1$  and  $T_1$  states (Fig. S3, ESI<sup>†</sup>), the broad

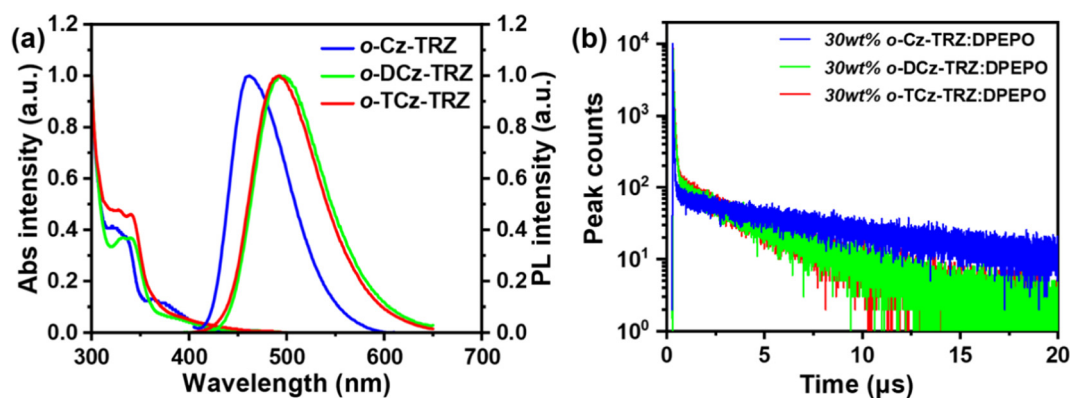


Fig. 2 (a) UV-vis absorption and PL spectra of *o*-Cz-TRZ, *o*-DCz-TRZ and *o*-TCz-TRZ in diluted toluene solution. (b) Transient PL decay curves of *o*-Cz-TRZ, *o*-DCz-TRZ and *o*-TCz-TRZ doped in DPEPO films (30 wt%).

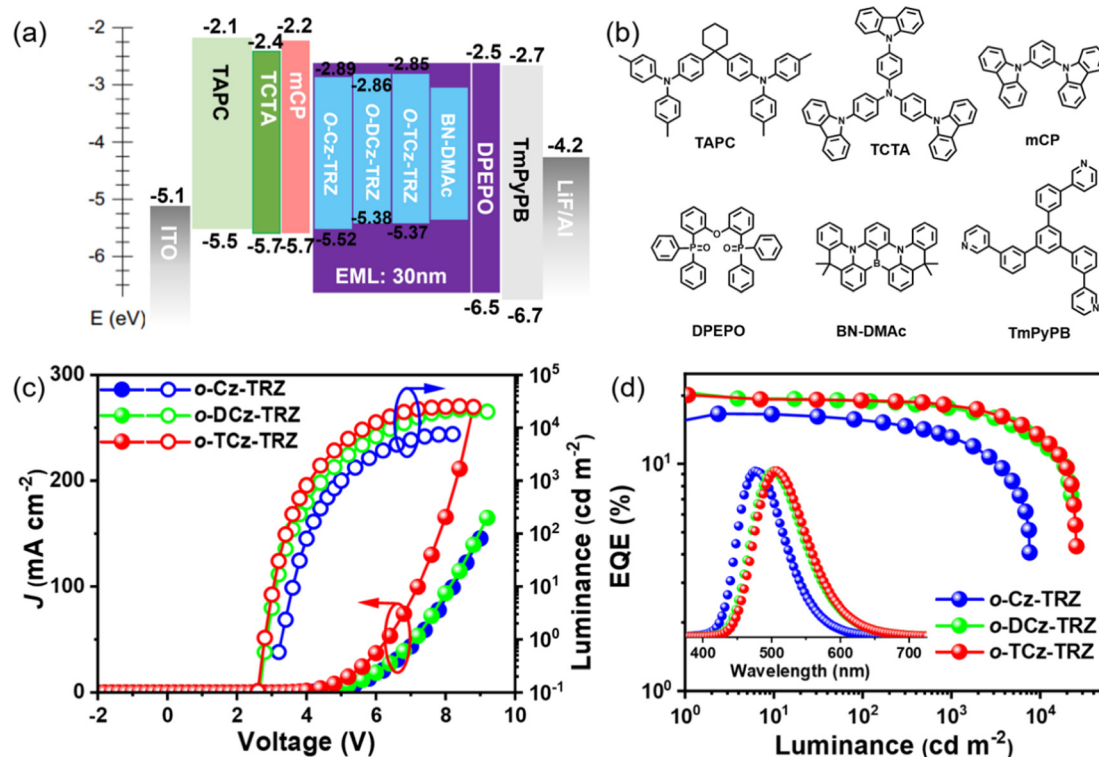


Fig. 3 (a) Device architecture and its energy level diagram. (b) Chemical structures of the corresponding materials. (c) Current density–Voltage–Luminance ( $J$ – $V$ – $L$ ) curves, (d) EQE–luminance curves and EL spectra (inset) of the devices based on *o*-Cz-TRZ, *o*-DCz-TRZ and *o*-TCz-TRZ as the emitter.

Table 1 Photophysical properties of the investigated molecules doped into DPEPO films at the concentration of 30 wt% at room temperature

Compound	$\lambda_{em}$ (nm)	PLQY (%)	$\phi_{PF}$ (%)	$\phi_{DF}$ (%)	$\phi_{ISC}$ (%)	$\tau_{PF}$ (ns)	$\tau_{DF}^a$ ( $\mu$ s)	$k_F$ ( $10^6$ s $^{-1}$ )	$k_{IC}$ ( $10^6$ s $^{-1}$ )	$k_{ISC}$ ( $10^7$ s $^{-1}$ )	$k_{RISC}$ ( $10^5$ s $^{-1}$ )	$k_F/k_{IC}$	$k_{RISC}/k_{ISC}$ ( $10^{-2}$ )
<i>o</i> -Cz-TRZ	475	65.7	26.3	39.4	73.7	26.7	8.2	9.8	5.1	2.8	2.5	1.9	0.9
<i>o</i> -DCz-TRZ	495	82.5	24.3	51.5	75.7	58.9	3.8	4.1	1.3	1.3	7.3	3.2	5.6
<i>o</i> -TCz-TRZ	498	80.6	15.9	36.4	84.1	52.3	3.9	3.0	2.8	1.6	7.0	1.1	4.4

<sup>a</sup> Average lifetime calculated by  $\tau_{av} = \sum_i^A \tau_i^2 / \sum_i^A \tau_i$ , where  $A_i$  is the pre-exponential for lifetime  $\tau_i$ .

and featureless fluorescence and phosphorescence spectra confirm the CT characteristics of the singlet and triplet states. Significant positive solvatochromism effects were observed as the embodiment of the strong CT characteristics for all the materials (Fig. S6, ESI $^\dagger$ ).

The solid-state photophysical properties of all the compounds were investigated by doping them into a typical host material bis[2-(diphenylphosphino)phenyl] ether oxide (DPEPO)<sup>43</sup> in a concentration of 30 wt%. All these doped films exhibit the typical emissions from *o*-Cz-TRZ, *o*-DCz-TRZ and *o*-TCz-TRZ with emission peaks of 475, 495 and 498 nm, respectively (Fig. S8, ESI $^\dagger$ ). The energy levels of the  $S_1/T_1$  states in the 30 wt% doped films were estimated to be 2.92/2.85, 2.87/2.83 and 2.85/2.82 eV from the onset of the low-temperature fluorescence and phosphorescence spectra (Fig. S7, ESI $^\dagger$ ), resulting in tiny  $\Delta E_{ST}$ s of 70, 40 and 30 meV for *o*-Cz-TRZ, *o*-DCz-TRZ and *o*-TCz-TRZ, respectively, which are conducive to the RISC process and TADF emission.

To verify the TADF characteristics, transient PL decay curves were measured in the doped films and are provided in Fig. 2b and Fig. S7 (ESI $^\dagger$ ). All these materials exhibit double-exponential decays with prompt lifetime ( $\tau_{PF}$ ) of 26–59 ns and delayed lifetime ( $\tau_{DF}$ ) of 3.8–8.2  $\mu$ s, demonstrating their TADF characteristics, which is in agreement with the tiny experimental  $\Delta E_{ST}$ s. Furthermore, from the temperature-dependent transient PL decay curves, the enhancement of the delayed fluorescence ratio with increased temperature from 100 K to 300 K further confirms the TADF characteristics (Fig. S7, ESI $^\dagger$ ).<sup>1–3</sup> The PLQYs of the *o*-Cz-TRZ, *o*-DCz-TRZ and *o*-TCz-TRZ doped films were measured to be 65.7%, 82.5% and 80.6%, respectively. Based on the transient PL decay spectra in the microsecond scale and PLQYs, high  $k_F$  of  $2.0$ – $9.8 \times 10^6$  s $^{-1}$  and  $k_{RISC}$  of  $2.5$ – $7.3 \times 10^5$  s $^{-1}$  were simultaneously achieved for these molecules (Table 1). As expected,  $k_F$  and  $k_{RISC}$  could be modulated by simply changing the number of carbazole units, which confirms the effectiveness of the molecular

design strategy, *i.e.*, modulating the PL properties and the exciton dynamic processes through adjusting the balance between intramolecular TBCT and TSCT. With increasing number of carbazole units, the  $k_F$  diminishes gradually mainly due to the gradually reduced overlap of the HOMOs and LUMOs. It follows that both intramolecular TBCT and TSCT contribute to promoting  $k_F$  but TBCT through the benzene linker (overlap of HOMOs and LUMOs locate at the linker) plays a more significant role than TSCT. Combining  $k_F$  and  $k_{IC}$  and comparing their ratios, that is,  $k_F/k_{IC}$  are 1.9, 3.2 and 1.1 for *o*-Cz-TRZ, *o*-DCz-TRZ and *o*-TCz-TRZ, respectively, the variation of PLQY (65.7%, 77.9% and 52.3% for *o*-Cz-TRZ, *o*-DCz-TRZ and *o*-TCz-TRZ, respectively) could be explained, which demonstrates the effectiveness of modulating  $k_F$  and PLQY by simply manipulating the proportion between intramolecular TBCT and TSCT. The  $k_{RISC}$ s of *o*-DCz-TRZ ( $7.3 \times 10^5 \text{ s}^{-1}$ ) and *o*-TCz-TRZ ( $7.0 \times 10^5 \text{ s}^{-1}$ ) are higher than that of *o*-Cz-TRZ ( $2.5 \times 10^5 \text{ s}^{-1}$ ), which can mainly be attributed to the stronger non-covalent interactions contributing to the strengthened SOC and the lower  $\Delta E_{ST}$ . As shown in Table 1, the  $k_{RISC}/k_{ISC}$  values are 0.9, 5.6 and  $4.4 (\times 10^{-2})$  for *o*-Cz-TRZ, *o*-DCz-TRZ and *o*-TCz-TRZ, respectively, demonstrating that the RISC processes are effectively promoted by greater intramolecular TSCT characters, which contributes to a higher triplet exciton utilization. Furthermore, the higher triplet exciton utilization and the higher  $k_{RISC}$  are conducive to high EQE and reduced efficiency roll-off especially at a high current density for a device.

From the above results and analysis, it is not difficult to find that TBCT plays a more significant role in promoting  $k_F$  *via* the overlap of the HOMOs and LUMOs at the benzene linker while intramolecular TSCT tends to contribute more to modulating  $k_{RISC}$  through the non-covalent weak interactions. Besides, it is worth noting that  $k_{RISC}$  can be improved with a relatively higher proportion of intramolecular TSCT characteristic but excessive intramolecular TSCT components have no significant effects on further promotion of  $k_{RISC}$  mainly on account of the reduced TBCT components, which also contribute to the promoted  $k_{RISC}$ . Therefore, the balance between intramolecular TSCT and TBCT for achieving relatively balanced  $k_{RISC}$ ,  $k_F$  and PLQY should be clarified and emphasized. An optimum balance could be found for *o*-DCz-TRZ, which possesses the highest PLQY and  $k_{RISC}$  and a relatively high  $k_F$ , making it a potential candidate to achieve high EQE and low efficiency roll-off in the corresponding device.

To evaluate the performances of those materials as assistant hosts, a green MR-TADF emitter BN-DMAC with an emission peak at 484 nm, a small FWHM of 29 nm and a high PLQY of 88% was used as the terminal emitter.<sup>29,30</sup> Significant absorption–emission spectral overlap was found between BN-DMAC and the assistant hosts (Fig. S10, ESI<sup>†</sup>), indicating the efficient energy transfer from the assistant hosts to BN-DMAC. The emission spectra of the ternary doped films DPEPO: 30 wt% sensitizers: 1 wt% BN-DMAC were measured and illustrated in Fig. S11 (ESI<sup>†</sup>). In all cases, the emission from the assistant hosts is almost eliminated with a peak at 494 nm and slightly broadened FWHMs of 40–43 nm, indicating the efficient energy transfer from the assistant hosts to BN-DMAC. The slightly

broadened FWHMs of these films can be attributed to the high polarity of DPEPO and these assistant hosts. In addition,  $\tau_{DF}$ s of 19.8–21.9  $\mu\text{s}$  were detected while using these molecules as the assistant host. Compared with BN-DMAC doped in the conventional DPEPO host, increased proportions of the delayed components were realized (Fig. S11 and Table S2, ESI<sup>†</sup>), indicating that the device performance of BN-DMAC can be improved by the assistant hosts under these conditions.

### OLED performance

To verify the application potential of these compounds and understand the relationship between the electroluminescence (EL) performances and the molecular structures, OLED devices were fabricated with a device architecture of ITO/TAPC (30 nm)/TCTA (10 nm)/mCP (10 nm)/DPEPO: 30 wt% emitters (30 nm)/DPEPO (10 nm)/TmPyPB (40 nm)/LiF (1 nm)/Al (150 nm), where (1,1-bis(4-(*N,N*-di(*p*-tolyl)-amino)-phenyl)cyclohexane) (TAPC), 4,4',4''tris(*N*-carbazolyl)triphenylamine (TCTA), 1,3-bis(*N*-carbazolyl)benzene (mCP), DPEPO, 1,3,5-tri(*m*-pyrid-3-ylphenyl)benzene (TmPyPB) and LiF layers play the roles of hole injection and hole-transport, electron-blocking, hole-blocking, electron transport and electron injection layers, respectively, as illustrated in Fig. 3. The device characteristics are depicted in Fig. 3 and Fig. S12 (ESI<sup>†</sup>) and the EL parameters are summarized in Table S4 (ESI<sup>†</sup>).

The devices based on *o*-Cz-TRZs, *o*-DCz-TRZ and *o*-TCz-TRZ as the emitter exhibit sky-blue to green emissions with peaks of 477, 500 and 505 nm, and the corresponding Commission Internationale de l'Eclairage (CIE) color coordinates are (0.16, 0.27), (0.23, 0.46) and (0.24, 0.49), respectively (Fig. 3d and Table S3, ESI<sup>†</sup>). Noticeably, *o*-DCz-TRZ and *o*-TCz-TRZ exhibit higher efficiencies with maximum EQEs of 20.3% and 20.1%, maximum power efficiencies (PEs) of 64.9 and 64.8  $\text{lm W}^{-1}$ , and maximum current efficiencies (CEs) of 57.8 and 57.8  $\text{cd A}^{-1}$ , respectively, owing to their higher PLQYs. Furthermore, benefitting from the high  $k_{RISC}$ , low efficiency roll-offs of 11.8% and 9.9% were also observed for the devices based on *o*-DCz-TRZ and *o*-TCz-TRZ (Fig. 3 and Table S3, ESI<sup>†</sup>) at the practical luminance of 1000  $\text{cd m}^{-2}$ , respectively. The simulated exciton annihilation processes of all the devices based on triplet–triplet annihilation (TTA) and singlet-polaron annihilation (SPA) models agree well with the experimental results, as shown in Fig. S13 (ESI<sup>†</sup>),<sup>44–46</sup> which demonstrates that TTA is the dominant mechanism for exciton annihilation at low current density and SPA can account for the efficiency roll-off at high current density in these devices.

The devices using these materials as the assistant host and BN-DMAC as the terminal emitter were fabricated in a structure of ITO/TAPC (30 nm)/TCTA (10 nm)/mCP (10 nm)/DPEPO: 30 wt% sensitizers: 1 wt% BN-DMAC (30 nm)/DPEPO (10 nm)/TmPyPB (40 nm)/LiF (1 nm)/Al (150 nm). In the ternary doped emission layer, the concentration of the terminal emitter BN-DMAC was optimized to 1 wt% to reduce the Dexter energy transfer process from the hosts. For comparison, the device without an assistant host was also fabricated. The devices with assistant hosts exhibit significantly suppressed efficiency roll-off and improved device performance owing to the high  $k_r$  of

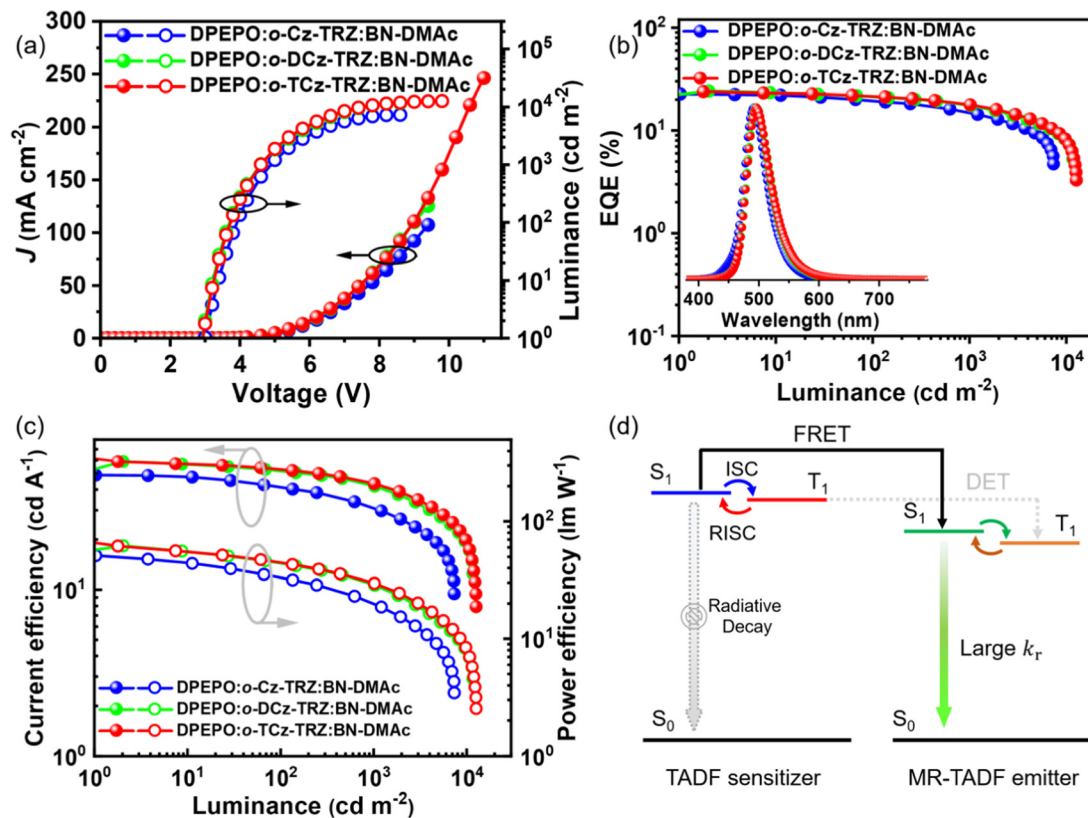


Fig. 4  $J$ - $V$ - $L$  curves (a), efficiency curves (b and c) and energy transfer mechanism (d) of the devices using *o*-Cz-TRZ, *o*-DCz-TRZ and *o*-TCz-TRZ as assistant hosts.

BN-DMAc and high  $k_{\text{RISC}}$  of the sensitizers. The Förster energy transfer (FRET) from the assistant host to the dopant can be quantified by utilizing the rate of FRET ( $k_{\text{ET}}$ ), the efficiency of FRET ( $\phi_{\text{ET}}$ ) and the FRET radius ( $R_0$ ). The results show that *o*-Cz-TRZ exhibits small  $R_0$  and lower  $k_{\text{ET}}$  and  $\phi_{\text{ET}}$  than those of *o*-DCz-TRZ and *o*-TCz-TRZ (Table S4, ESI<sup>†</sup>), evidencing the more efficient energy transfer from *o*-DCz-TRZ and *o*-TCz-TRZ to BN-DMAc. Fig. 4b depicts the EL spectra of the *o*-Cz-TRZ, *o*-DCz-TRZ and *o*-TCz-TRZ sensitized devices, exhibiting FWHMs of 44, 46 and 46 nm with identical emission peaks of 492, 496 and 496 nm and corresponding CIE coordinates of (0.12, 0.40), (0.13, 0.49) and (0.14, 0.49), respectively. The  $V_{\text{on}}$ s of these devices are 2.9–3.0 V (Fig. 4a), indicating efficient carrier injection and transport characteristics. Similarly, the devices using *o*-DCz-TRZ and *o*-TCz-TRZ as assistant hosts achieve better performance, with maximum EQEs of 24.2% and 23.9%, CEs of 59.1 and 61.7 cd A<sup>-1</sup> and PEs of 61.9 and 66.0 lm W<sup>-1</sup> (Fig. 4 and Table S5, ESI<sup>†</sup>), respectively, which are higher than those of the devices with the same terminal emitter BN-DMAc using mCBP or DPEPO hosts (Fig. S14 and Table S5, ESI<sup>†</sup>) and comparable to the device using mCBP and PO-T2T exciplex hosts (Table S6, ESI<sup>†</sup>). What's more, compared with the DPEPO hosted device (Fig. S14 and Table S5, ESI<sup>†</sup>), these devices achieve higher efficiency and luminance and reduced efficiency roll-off. For example, the *o*-TCz-TRZ sensitized device exhibits EQEs of 21.5% and 17.9% at the practical luminance of

100 and 1000 cd m<sup>-2</sup>, respectively, corresponding to efficiency reduction of 10.0% and 16.9% relative to the maximum EQE, which are also better than the devices using typical mCBP and exciplex hosts (mCBP:PO-T2T).<sup>29,30</sup> After the introduction of assistant hosts, the predominant energy transfer pathway is changed to the long-range FRET from the assistant host to BN-DMAc, the population of triplet excitons on BN-DMAc can be greatly reduced to improve the exciton utilization and reduce the corresponding quenching effect (Fig. 4d), resulting in high efficiency and reduced efficiency roll-off.

## Conclusion

In summary, three TADF molecules are successfully designed and synthesized through the *ortho*-linkage of the diphenyltriazine acceptor and the oligo carbazole donor to the benzene linker. Combining the computational simulation and experimental analysis, the concurrent intramolecular TBCT and TSCT are confirmed and the ratios are successfully modulated by simply controlling the number of carbazole units. Through delicately balanced intramolecular TSCT and TBCT, relatively high  $k_{\text{F}}$  and  $k_{\text{RISC}}$  are achieved simultaneously. Benefitting from the high PLQY and  $k_{\text{RISC}}$ , the device using *o*-DCz-TRZ as an emitter gives maximum efficiencies of 20.3% and 64.9 lm W<sup>-1</sup>, characterized by slow efficiency roll-off with EQE of 17.9% at

the luminance of 1000 cd m<sup>-2</sup>. Furthermore, by using these materials as assistant hosts and sensitizers and DMac-BN as a terminal emitter, high efficiencies of 23.9% and 66.0 lm W<sup>-1</sup> are achieved with FWHM of 46 nm, featuring slow efficiency roll-off with EQE of 17.9% at the luminance of 1000 cd m<sup>-2</sup>. This reveals the importance of modulating the balance of intramolecular TBCT and TSCT in TADF materials towards high  $k_F$  and  $k_{RISC}$  rates and thus high efficiency and low efficiency roll-off OLEDs.

## Conflicts of interest

There are no conflicts to declare.

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